This listing of claims replaces all prior versions and listings of claims in this application.

LISTING OF CLAIMS:

Claim 1 (Previously Presented): A tablet comprising crystals of a pharmaceutically acceptable salt of citalogram, wherein the median particle size of the crystals is at least 40 μ m, which is prepared by direct compression of the pharmaceutically acceptable salt and pharmaceutically acceptable excipients.

Claims 2-3 (Canceled)

Claim (Previously Presented): The tablet according to claim 1 which does not contain a binder.

Claim & (Previously Presented): The tablet according to claim 1 which contains 2-60% w/w active ingredient calculated as citalogram base.

Claim (Previously Presented): The tablet according to claim 1 which contains a filler selected from lactose, sugars, calcium phosphates, starch, modified starches, microcrystalline cellulose, calcium sulfate and calcium carbonate.

Claim / (Previously Presented): The tablet according to claim s, wherein the filler is a microcrystalline cellulose.

Docket No.: 542 2/01004 Page 2

Claim & (Previously Presented): The tablet according to claim 1 which contains a lubricant selected from metallic stearates, stearic acid, wax, hydrogenated vegetable oil, tilc and colloidal silica.

Claim (Previously Presented): The tablet according to claim 8, wherein the lubricant is magnesium stearate or calcium stearate.

Claim 19 (Previously Presented): The tablet according to claim 1 which is substantially free of lactose.

Claim 11 (Canceled)

Claim 12 (Previously Presented): The tablet according to claim 1 wherein the pharmaceutically acceptable salt is citalopram hydrobromide or citalopram hydrochloride.

Claim 1/3 (Previously Presented): The tablet according to claim 1/2, wherein the pharmaceutically acceptable salt is citalopram hydrobromide.

Claims 14-35 (Canceled)

Claim 36 (Previously Presented): The tablet of claim 1, which contains 10-40% w/w active ingredient calculated as citalogram base.

Claim 3/7 (Previously Presented): The tablet of claim 1, which contains 15-25%

w/w active ingredient calculated as citalopram base.

U.S. Patent Application Serial No. 09/730,380

A mendment

Docket No.: 542 2/01004 Page 3

09/18/03 THU 11:10 [TX/RX NO 5201] 2005

Claim 38 (Previously Presented): The tablet of claim 8, wherein said filler is a sugar selected from the group consisting of sorbitol, mannitol, dextrose and sucrose.

Claim 39 (Previously Presented): The tablet of claim 6, wherein said filler is a calcium phosphate selected from the group consisting of dibasic, tribasic, hydrous and anhydrous calcium phosphate.

Claim 40 (Previously Presented): The tablet of claim 8, wherein said lubricant is a metallic stearate selected from the group consisting of magnesium, calcium and odium stearate.

Claim 4 (Previously Presented): The tablet of claim 1, wherein the crystals have a median particle size of 40-200 μ m.

Claim 42 (Previously Presented): The tablet of claim 1, wherein the crystals have a median particle size of 45-150 \mu m.

Claim 43 (Previously Presented): The tablet of claim 1, wherein the crystals have a median particle size of 50-100 µm.

Claims 44-92 (Canceled)

Docket No.: 542 2/01004 Page 4